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Thickness Dependence of Magneto-transport Properties in Tungsten Ditelluride

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We investigate the electronic structure of tungsten ditelluride (WTe_2) flakes with different thicknesses in magneto-transport studies. The temperature-dependent resistance and magnetoresistance (MR) measurements both confirm the breaking of carrier balance induced by thickness reduction, which suppresses the ‘turn-on’ behavior and large positive MR. The Shubnikov-de-Haas oscillation studies further confirm the thickness-dependent change of electronic structure of WTe_2 and reveal a possible temperature-sensitive electronic structure change. Finally, we report the thickness-dependent anisotropy of the Fermi surface, which reveals that multi-layer WTe_2 is an electronic 3D material and the anisotropy decreases as thickness decreases.

I. INTRODUCTION

Tungsten ditelluride (WTe_2), a layered transition metal dichalcogenide (TMD) material, has attracted a great deal of interest due to its unique electronic transport properties since the discovery of the non-saturating positive large magnetoresistance (MR) in bulk¹. It is widely believed that the extraordinary MR comes from the nearly perfect balance between electron and hole concentrations¹⁻⁷. Many other peculiar electronic properties have also been observed in WTe_2 in transport measurements, such as ‘turn-on’ behavior^{1,8-10}, multi-Fermi pockets revealed by Shubnikov-de-Haas (SdH) oscillations¹¹⁻¹³, surprisingly small Fermi surface anisotropy⁴, ferroelectricity¹⁴⁻¹⁷, superconductivity¹⁸⁻²⁰, etc. WTe_2 is also a topological material. In the bulk form, WTe_2 has been predicted²¹ and observed²² to be a type-II Weyl semimetal. In the monolayer form, WTe_2 is a quantum spin Hall insulator^{23,24} at low carrier density (n). However, it is still unclear how the topological property and the electronic structure evolve by reducing the crystal thickness.

Here we investigate the evolution of electronic structure in WTe_2 flakes with different thicknesses by performing temperature dependent resistance (R - T) measurements, MR measurements, SdH oscillation studies and angle-dependent MR measurements. Experiments show that the imbalance of carrier densities caused by thickness reduction plays an important role in the suppression of the ‘turn-on’ behavior and the large positive MR, while the non-saturating characteristic was hardly affected. We further confirm that the multi-layer WTe_2 is also an electronic 3D material like the bulk crystal and that the anisotropy reduces as thickness decreases.

II. EXPERIMENT

WTe_2 flakes with different thicknesses were obtained by mechanical exfoliation of bulk WTe_2 crystals synthesized by chemical vapor transport⁹. Six exfoliated flakes characterized in this paper can be classified into three groups. The first group contains two thick samples (denoted as ‘sample 1’ and ‘sample 2’ hereafter) with thickness around 150 nm, the second group contains three thin flakes with thickness around 20 nm (denoted as ‘sample 3’ through ‘sample 5’ hereafter), and the third group contains one ultra-thin flake with thickness at 5 nm (‘sample 6’), whose transport properties have been reported elsewhere²⁵. In this paper, we will focus on the first two groups. The thick flakes are directly transferred onto silicon substrates with 285 nm-thick SiO_2 coating on the surface. The thin flakes were encapsulated between two pieces of hexagonal boron nitride (hBN) thin flakes which were about 10 nm thick and transferred onto the substrates by a dry transfer technique²⁶. Thin WTe_2 flakes get oxidized easily upon exposure to air^{27,28}. Hence the hBN flakes are necessary here to protect the thin samples from air-induced degradation. In addition, hBN flakes provide a cleaner interface for WTe_2 . For the thick samples, photo-lithography was used to make the patterns. For the thin samples, electron-beam lithography was used to make patterns. The Ohmic contacts were deposited by electron-beam evaporation of Pd/Au (10 nm/200 nm for thick samples, 10 nm/40 nm for thin samples) followed by a lift-off process. Transport measurements down to 0.02 K were carried out in an Oxford dilution refrigerator. Both the longitudinal resistance R_{xx} and Hall resistance R_{xy} were measured simultaneously by using standard low frequency lock-in techniques.

III. R - T CHARACTERISTICS AND ‘TURN-ON’ BEHAVIOR

Except in the ultra-thin sample, the temperature-dependent longitudinal resistances (R - T) in all other five devices (samples 1-5) with different thicknesses show metallic properties across the full experiment temperature range, as shown in FIG. 1(a). The ultra-thin sample (sample 6), however, shows an insulating temperature dependence at low temperature region and it is more than 10 times more resistive than all other samples. All samples in the same group show similar behavior. Thus, we pick up sample 2 and sample 3 as representatives for the thick and thin groups, respectively. In FIG. 1(b), the R - T curves for sample 2 and sample 3 are plotted in log-log scale. Fermi liquid fits, $R_0 = \alpha + \beta T^2$, applied at low temperatures are also shown as the black dashed lines for both samples. Here R_0 represents the resistance at 0 T magnetic field and α, β are two fitting parameters. We found that the maximum applicable T of the Fermi liquid fit decreased from ~ 60 K in sample 2 to ~ 40 K in sample 3 with decreasing thickness. Comparing with the case in bulk (~ 80 K)⁸, the trend shows consistently that the applicable T range decreases with thickness decreasing.

We further investigated the R - T curves at various magnetic fields up to 12 T. We unambiguously observed the ‘turn-on’ behavior in sample 2, as shown in FIG. 1(c), in which $(R(T) - R(30 \text{ K}))/R(30 \text{ K})$ is plotted as a function of T below 30 K. The R - T curves gradually change from metallic to insulating with magnetic field increasing from 0 T to 12 T. The critical field $\mu_0 H^*$ is around 7 T, which is much larger than the one reported in bulk (below 2 T)^{1,8,9}. Due to the fact that the ‘turn-on’ behavior only occurs in the Fermi liquid state^{8,29}, we assume that the larger H^* in our case is caused by the shift of the Fermi liquid state to a lower temperature region due to the thickness decreasing. Our assumption can be supported by a comparison measurement in sample 3. In sample 3, the ‘turn-on’ behavior can’t be observed up to 12 T. In a higher field, SdH oscillations emerge and disguise the ‘turn-on’ behavior. That means a magnetic field larger than 12 T is required to manifest the ‘turn-on’ behavior with the Fermi liquid state further moving to a lower temperature region (below 40 K).

In order to investigate the origin of the ‘turn-on’ behavior, we applied Kohler’s rule:

$$\text{MR} = A(H/R_0)^m \quad (1)$$

in which R_0 is the resistance of R - T curve at zero magnetic field and A, m are the constant fitting parameters. Here MR is defined as $\text{MR} = [R(T, H) - R_0(T)]/R_0(T)$. We found that the R - T curves in FIG. 1(c) can be scaled into one curve. Specifically, the Kohler’s rule fitting gave the parameter $m = 1.8$. Comparing with the case in bulk ($m = 1.92$)⁸, m decreased and deviated further from the perfect carrier compensation of $m = 2$ ^{30–32}. The scaling behavior obtained by Kohler’s rule confirmed that the R - T curves at different magnetic fields have the same temperature dependence, although it looks like the curve at higher field shows larger MR effect. This is quite different from the case of a magnetic field-induced metal-insulator transition, which requires a larger increasing rate at a higher magnetic field due to gap opening^{29,33}. In addition, our results show that the ‘turn-on’ behavior takes its origin from carrier compensation. The nearly broken carrier compensation in sample 3 ($m = 1.69$) makes the ‘turn-on’ behavior almost invisible even at 12 T.

IV. TWO-BAND MODEL FITTINGS

To further confirm whether perfect carrier compensation is related to the ‘turn-on’ behavior or not, we performed MR measurements on both samples. The MR curves of both samples at different temperatures are shown in FIG. 2(a) and 2(d). We found that the MR curves in sample 2 crossed at two points located at 7 T and -7 T, respectively. This is consistent with the $\mu_0 H^*$ of the ‘turn-on’ behavior. However, it’s almost impossible to identify such two crossing points in sample 3 since the MR curves at different temperatures are fully overlapped in large magnetic field region. This is also consistent with the fact that we didn’t observe obvious ‘turn-on’ behavior in sample 3.

Quasi-quadratic positive MR can be seen in both samples, which has been attributed to the perfectly balanced electron and hole densities^{3–7,9}. Comparing with the results in bulk¹, which recorded MR as high as 13,000,000 %, the highest MR in our samples only showed 1,200 % due to reduction in thickness. The MR in both samples is notably suppressed, which indicates an imperfect balance between carrier densities and more impact from defects. But there’s still no observed trend towards saturation of MR in the samples. In order to examine the balance between carrier densities further, we additionally performed Hall measurements. Combined with MRs, we could extract the carrier densities and mobilities using the two-band model^{34–36}:

$$R_{xx} = \frac{\sigma_1 + \sigma_2 + (\sigma_1 \sigma_2^2 R_{H_2}^2 + \sigma_2 \sigma_1^2 R_{H_1}^2) B^2}{(\sigma_1 + \sigma_2)^2 + \sigma_1^2 \sigma_2^2 (R_{H_1} + R_{H_2})^2 B^2} \quad (2)$$

$$R_{xy} = \frac{R_{H_1} \sigma_1^2 + R_{H_2} \sigma_2^2 + \sigma_1^2 \sigma_2^2 R_{H_1} R_{H_2} (R_{H_1} + R_{H_2}) B^2}{(\sigma_1 + \sigma_2)^2 + \sigma_1^2 \sigma_2^2 (R_{H_1} + R_{H_2})^2 B^2} \quad (3)$$

in which $\sigma_i = n_i e \mu_i$ ($i = 1, 2$) are the conductance contributions from electron and hole, respectively. $R_{H_i} = 1/n_i e$ ($i = 1, 2$) are the Hall coefficients for electron-dominant and hole-dominant Hall effect. n_i and μ_i are carrier densities and mobilities, respectively. The fitting results and the extracted carrier densities and mobilities are shown in FIG. 2.

The two-band model was perfectly applicable to our MR and Hall measurements for sample 2, while there’s a little mismatch in the low field region of MR for sample 3 (Fig. 2(e)). The low-field mismatch might come from the disorder-induced quantum interference effect^{25,37–39}, which is more pronounced in ultra-thin sample where weak localization and weak anti-localization effects are stronger. However, this small mismatch doesn’t have decisive influence on the extraction of carrier densities and mobilities. In sample 2, we found that the hole density was almost unchanged with temperature while the electron density gradually increases with temperature increasing (Fig. 2(c)). Such a trend is consistent with other transport and ARPES studies^{3–7}. However, the electron and hole densities are not completely equal at low temperature in our samples. The charge carrier density ratio, n_e/n_h , is about 1.15 in sample 2 and becomes even larger in sample 3, which it is 1.26. It indicates that the perfect balance between electron and hole carrier densities will be gradually broken as the thickness decreases. Furthermore, such a growing charge imbalance will cause the MR to be continuously suppressed and will also cause the ‘turn-on’ behavior to become insignificant and eventually disappear. However, it seems that the MR can still maintain the non-saturating characteristic at large field up to 12 T, no matter how much the MR is suppressed. While the MR in some imbalanced-carrier systems tends to saturate eventually at high magnetic fields^{40,41}, it is not observed in our experiments. This could be attributed to the fact the either the magnetic field used in our experiment is not large enough or the origin of the non-saturating characteristic is not the perfect balance of electron and hole compensation. Actually a linear non-saturating MR has been observed in some Dirac semimetals^{42–46}, which is believed to take the origin of the lifting of a remarkable protection mechanism induced by time reversal symmetry that strongly suppresses backscatterings at zero magnetic fields. Similar linear MR curves have been observed in disordered WTe₂ flakes^{25,47}. A possible link between the non-saturating MR in WTe₂ and such a mechanism also deserves further investigation.

It is worth mentioning that both the electron and hole mobilities in sample 2 increased with decreasing temperature, which might be due to the suppression of phonon scattering at lower temperatures. In sample 3, the electron and hole mobilities were

larger than those in sample 2 and remained largely unchanged with temperature. This may indicate the mobility enhancement brought about by the encapsulating hBN flakes, which eliminate the interface scattering.

V. SHUBNIKOV DE HAAS OSCILLATIONS

SdH oscillations can be clearly seen in low temperature MR traces (FIG. 2(a) and 2(d)) due to the high carrier mobility and the strong suppression of phonon scattering at low temperatures. The SdH oscillations were extracted from the MR curves by removing the quasi-quadratic backgrounds. The extracted oscillations at different temperatures are shown in FIG. 3(a) and 3(d) for sample 2 and sample 3, respectively. FIG. 3(b) and 3(e) shows the corresponding Fourier transformation (FT) analysis. There were three obvious peaks observed in sample 2 and two peaks in sample 3. In sample 2, the three peaks are located at 102 T (α), 161 T (β) and 187 T (γ). These peak locations almost stay unchanged with temperature. In sample 3, one peak located at 113 T and the other one located at 176 T at 0.02 K and re-located to 169 T above 0.02 K. According to the similarity of locations in sample 2, we assigned α to the first peak and β to the second peak.

According to the Onsager relation $F = (\Phi_0/2\pi^2)A_F$, where F is the oscillation frequency, Φ_0 is the flux quantum, the cross-sectional area of each Fermi pocket can be obtained: $A_F = 0.0097, 0.0153, 0.0178 \text{ \AA}^{-2}$ for α, β, γ pockets in sample 2, respectively. And $A_F = 0.0108, 0.0168$ (at 0.02 K) and 0.0159 (above 0.02 K) \AA^{-2} for α and β pockets in sample 3, respectively. It is inaccurate to identify the carrier types of the Fermi pockets based on both the size of each Fermi pocket and the carrier densities obtained from the two-band model fittings, but we can conclude that the electronic structure changes dramatically with thickness. Firstly, the size of the corresponding pockets in the two samples is marginally different. Although in other SdH oscillation studies in WTe_2 , the size of the pockets is not the same due to different sample sources. In our case, sample 2 and 3 are from the same crystal. So such a difference in the size of the Fermi pockets can be attributed to the changes in the Fermi surface topology caused by the finite-size effect¹³. Secondly, one pocket (γ) totally disappears in sample 3. Compared with the bulk, in which four pockets are discovered¹¹⁻¹³, and the gated trilayer sample, in which two pockets are observed⁴⁸, the cases in our samples are in an intermediate state and in line with the trend that the smaller the thickness, the fewer the number of pockets. To sum up the two points, the difference could be intrinsically attributed to the size effect caused by the thickness change of the sample, but other effects, like scatterings from impurities⁴⁹, strain effect^{50,51} cannot be ruled out. Thirdly, the shift of the β pocket with temperature in sample 3 reveals a temperature-sensitive electronic structure of multi-layer WTe_2 . This might be due to the Lifshitz transition which has been previously reported in ARPES studies^{3,5} and band structure calculations⁵² in WTe_2 . Under certain temperatures, the sizes of some carrier pockets dramatically change due to temperature-induced Fermi surface shift. Besides the Fermi pocket topology, a temperature-induced spin splitting can be explicitly observed in sample 3, as shown in FIG. 3(d). The oscillation peaks double-split with increasing temperature, which is contrary to the common belief that the spin splitting only occurs at low temperature and high magnetic field. What happened here might be due to the breaking of spin-orbit coupling (SOC) with increasing temperature. It has been proved that SOC can be strongly suppressed by temperature-related scatterings in WTe_2 ²⁵. At low temperature, the SOC is strong so that the applied magnetic field is not sufficient to support an observable Zeeman splitting. At higher temperature, the missing or non-dominant SOC enables the appearance of the Paschen-Back effect at moderate magnetic field⁵³⁻⁵⁵. We did not observe splitting in sample 2, which may be because in thicker samples, the scattering is usually weak due to less fabrication caused disorders, so the SOC is still strong enough and preventing a similar splitting.

The Lifshitz-Kosevich (LK) formula is commonly used to analyze the SdH oscillations⁵⁶. The damping factor shown below in the LK formula is used to describe the temperature dependence of the oscillation amplitude,

$$R_T = \frac{2\pi^2 k_B T m^* / \hbar e B}{\sinh(2\pi^2 k_B T m^* / \hbar e B)} \quad (4)$$

in which k_B is the Boltzmann constant and m^* is the effective mass. FIG. 3(c) and 3(f) show the fitting results for the Fermi pockets in sample 2 and sample 3, respectively. Since the amplitude of the γ peak in sample 2 and δ peak in sample 3 cannot be reliably extracted above 3.5 K and the β peak in sample 3 shifts location, we excluded them from the fitting analysis. The effective mass obtained from fittings are: $m_\alpha = 0.393 m_e$ and $m_\beta = 0.419 m_e$ for sample 2, and $m_\alpha = 0.301 m_e$ for sample 3, where m_e is the free electron mass. We notice that sample 2 shows comparable results with those in bulk, whereas for sample 3, the effective mass is slightly lighter. This might result from the enhanced carrier mobilities due to the encapsulated structure^{11,13,57}. In addition, it is worth mentioning that, the mean free path of each sample can be estimated as $l_2 \approx 39 \text{ nm}$ in sample 2 and $l_3 \approx 46 \text{ nm}$ in sample 3 according to $l = \hbar k_F \mu / e$, in which k_F is the Fermi wave vector expressed as $k_F = \sqrt{A_F / \pi}$. This means that, to a certain extent, the sample quality has not deteriorated as the thickness of the sample decreases. The impurity, scattering effect, etc. in these two samples basically maintain the same level.

VI. FERMI SURFACE ANISOTROPY

We further measured the angle-dependent MR curves, as shown in FIG. 4(a) and 4(c) for sample 2 and sample 3, respectively. θ is the angle between the direction of the magnetic field and the normal direction of the sample, as shown in the inset of FIG. 4(b). It is observed that the non-saturating positive MR is strongly suppressed when the magnetic field is parallel to the sample surface. Magnetoresistance oscillation study shows that the peaks in the FT spectrum shift with angle, which indicates the Fermi surface anisotropy between the in-plane and out-of-plane directions. So we further investigated the anisotropy of the Fermi surface through the scaling behavior of MR curves. The scaling behavior^{4,58,59} can be expressed as:

$$R(H, \theta) = R(\varepsilon_\theta H) \quad (5)$$

with the scaling factor $\varepsilon_\theta = (\cos^{-2} \theta + \gamma^{-2} \sin^{-2} \theta)^{1/2}$. All the MR curves can be scaled into a single curve as shown in FIG. 4(b) and 4(d), respectively, for both samples. Since the sample resistance is directly related to the effective mass by $R = m^*/ne^2\tau$, where τ is the relaxation time, the scaling behavior of MR can describe the mass anisotropy m_{\parallel}/m_{\perp} , which is the γ in the equation. In addition, since m^* manifests the energy band curvature, γ alone can also be used to describe the anisotropy of Fermi surface k_{\parallel}/k_{\perp} ^{4,59-61}.

The scaling factors were obtained for both samples at different angles and plotted in FIG. 4(e). By fitting the scaling factor versus angle, we got the anisotropy parameter $\gamma = 8.08$ and 2.28 in sample 2 and sample 3, respectively. Interestingly, the anisotropy we got from WTe₂ is much smaller than that in layered material graphite (~ 12)⁵⁹, and that in 3D high-T_c superconductor YBa₂Cu₃O₇ (~ 9)⁶². Similar small anisotropy in WTe₂ has been experimentally confirmed in several other probes as well^{13,63-65}. It is even more striking that the anisotropy parameter in the thick sample is larger than the one in the thin sample. That means the electrons in a thinner WTe₂ sample can move more freely along k_z direction than in a thicker one. Such a small Fermi surface anisotropy indicates that WTe₂ is actually an electronic 3D material. Unlike the isotropic electron gas, however, the electron movement in WTe₂ along the stacking direction is strongly modulated by sample thickness. When the thickness reduces to dozens of nanometers, a thinner sample may suffer less interlayer scatterings along the stacking direction, thereby making the motion of electrons in this direction less constrained. Overall, the multi-layer WTe₂ is an electronic 3D material with even smaller anisotropy than that in bulk. This novel evolution of the Fermi surface anisotropy demands further investigations in few-layer or even monolayer materials.

VII. CONCLUSION

With a thorough magneto-transport study in a series of WTe₂ flakes, we have observed a systematic change of electronic structure as a function of the thickness. First, we confirmed that the Kohler's rule is applicable and responsible for the 'turn-on' behavior which normally occurs in Fermi liquid state, thereby ruling out the possibility of a metal-insulator transition. Second, we found that the imbalance of carrier densities took an important role on the suppression of 'turn-on' behavior and large positive MR, while the non-saturating characteristic was hardly affected. This might hint at some other origins for such a MR. Third, the SdH oscillation studies further shown the important role of thickness on the Fermi surface in WTe₂ and perhaps a temperature-sensitive change in electronic structure. Finally, we reported a thickness-dependent Fermi surface anisotropy, which revealed that WTe₂, a typical 2D Van der Waals material, is effectively an electronic 3D material and the anisotropy decreases with decreasing thickness.

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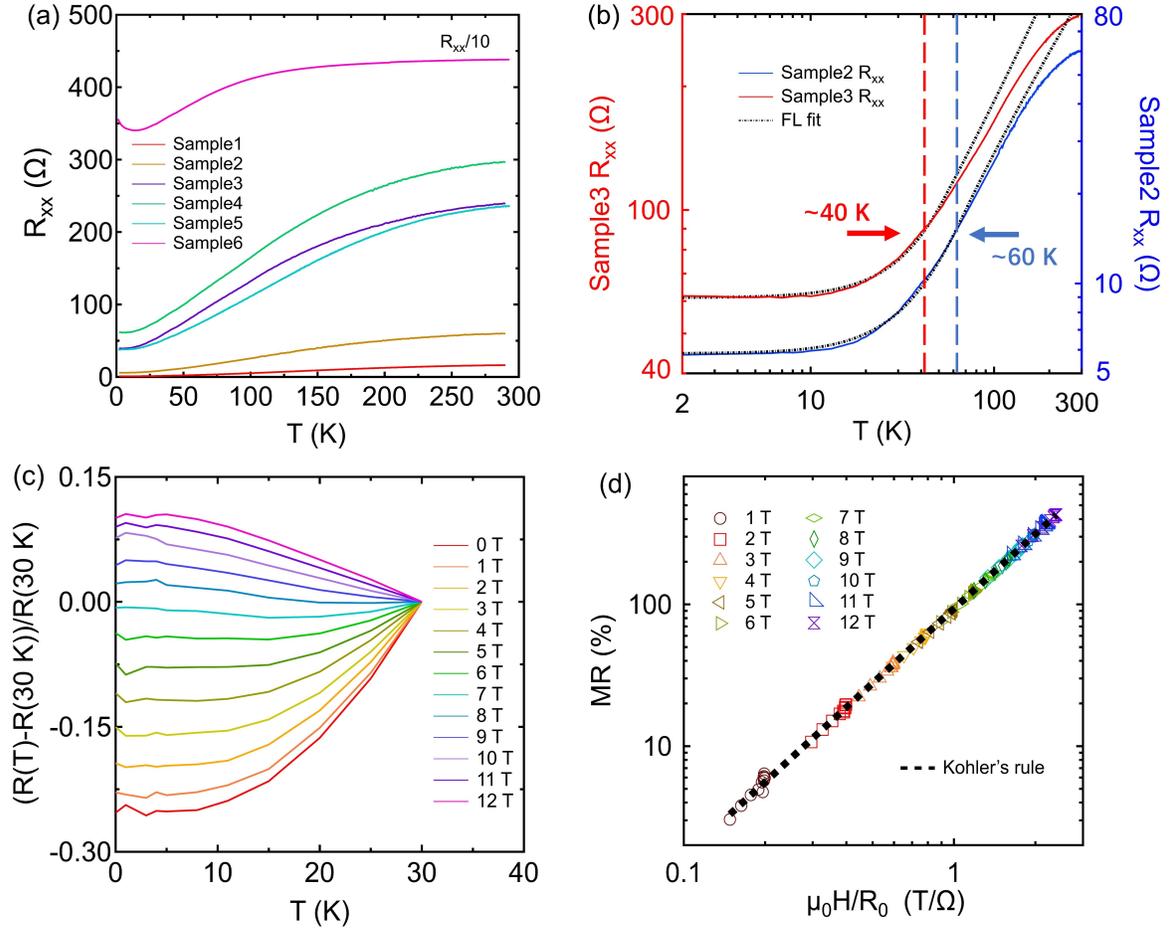


FIG. 1. Temperature-dependent resistance measurements. (a) R - T curves of six different devices. Five of them (samples 1-5) show metallic behavior. (b) Fermi liquid fits (dashed lines) for sample 2 and sample 3, respectively, at low temperature regions. The upper limits of the Fermi liquid regions are marked with colored dashed lines and arrows. (c) $(R(T) - R(30 \text{ K})) / R(30 \text{ K})$ curves at various magnetic fields manifest 'turn-on' behavior in sample 2. (d) Kohler's rule scaling of the data in panel (c).

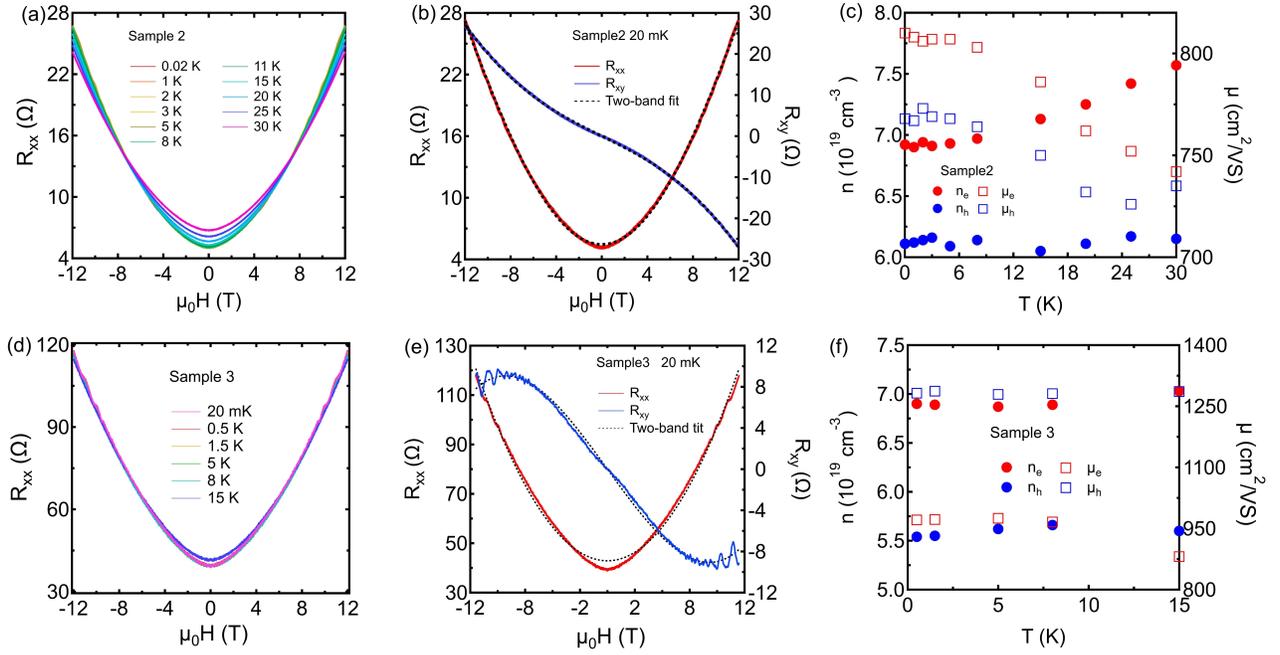


FIG. 2. Magnetoresistances and two-band model fittings in samples 2 and 3. (a) MR curves of sample 2 at various temperatures. (b) A representative two-band model fitting result for sample 2 at 20 mK. (c) Carrier densities (solid circles) and mobilities (open squares) for both electrons (red) and holes (blue) extracted from two-band model fittings at different temperatures for sample 2. (d) MR curves of sample 3 at various temperatures. (e) A representative two-band model fitting result for sample 3 at 20 mK. (f) Carrier densities (solid circles) and mobilities (open squares) for both electrons (red) and holes (blue) extracted from two-band model fittings at different temperatures for sample 3.

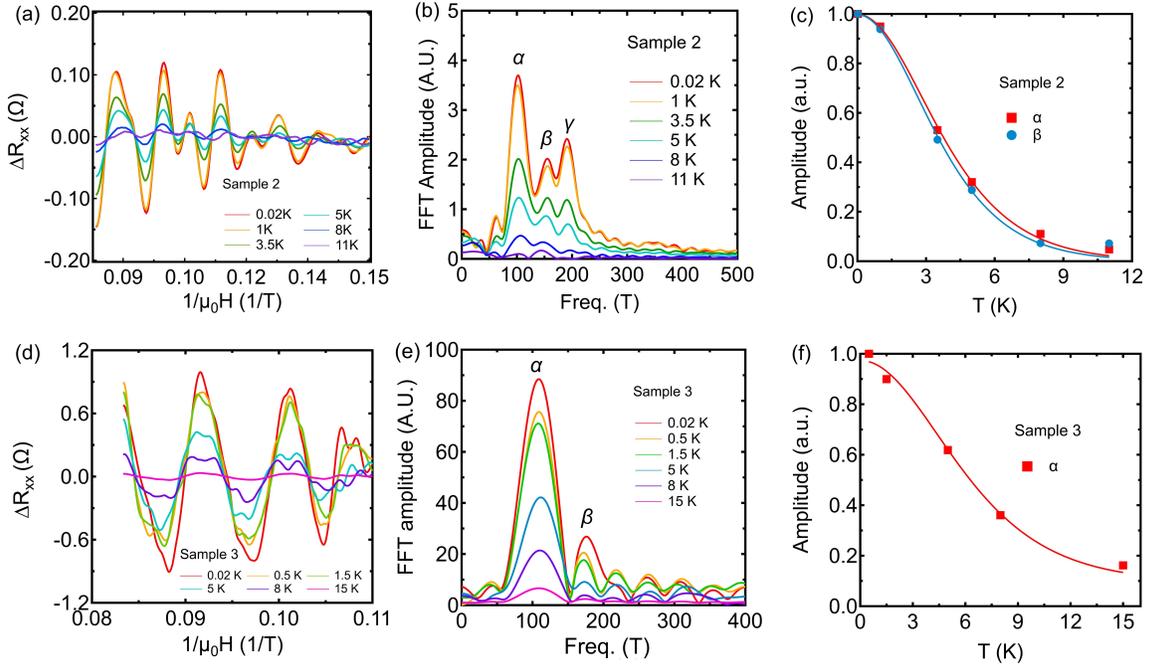


FIG. 3. Temperature dependence of SdH oscillations. (a) SdH oscillations extracted from the MR curves for sample 2. (b) FT analysis shows three peaks corresponding to three Fermi pockets in sample 2, which are marked as α , β and γ . (c) Temperature dependence of the amplitude of oscillation peaks (square and circle symbols) and the LK fit (solid lines) in sample 2. (d) SdH oscillations extracted from the MR curves for sample 3. (e) FT analysis shows two peaks corresponding to two Fermi pockets in sample 3, which are marked as α and β . (f) Temperature dependence of the amplitude of oscillation peaks (circle symbols) and the LK fit (solid lines) in sample 3.

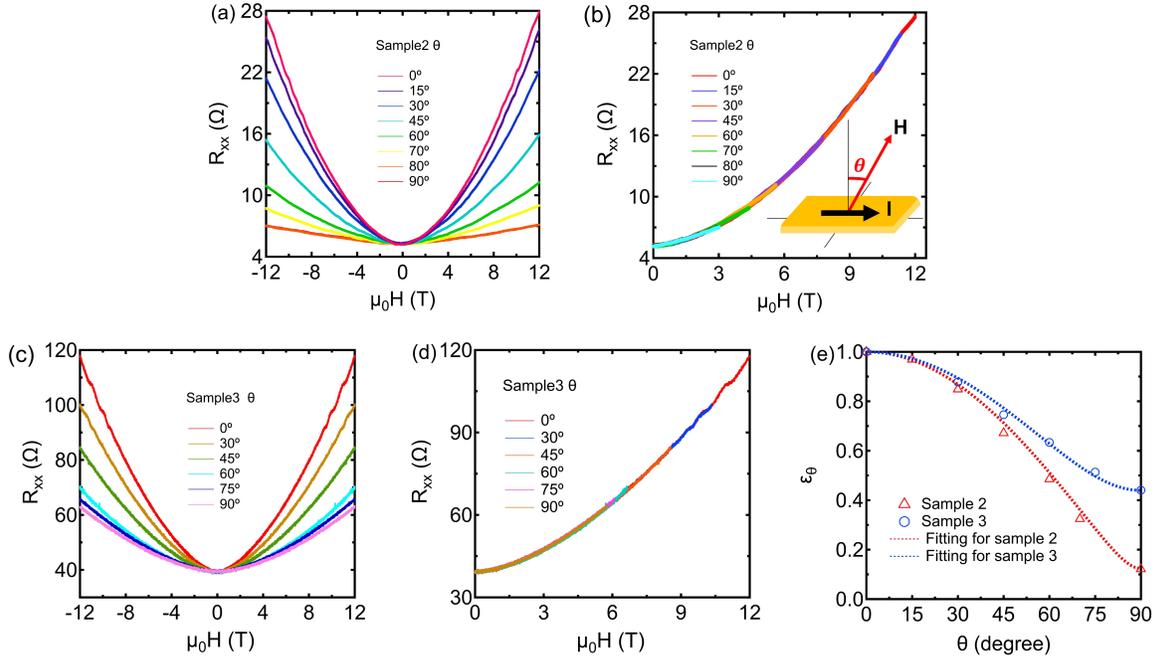


FIG. 4. Angle-dependent MR curves and the scaling behaviour. (a) Angle-dependent MR curves in sample 2. (b) The scaling behavior of the MR curves in sample 2. Inset shows the schematic of field orientation. (c) Angle-dependent MR curves in sample 3. (d) The scaling behavior of the MR curves in sample 3. (e) The scaling factors ϵ_θ at different angles (open symbols) and fittings (dotted lines) for both samples.